d his

(FILE 'HOME' ENTERED AT 11:17:00 ON 28 FEB 2007)

43 S L7 AND HEART?

17 S L16 AND PD<1999

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 11:21:33 ON 28 FEB 2007 20802 S LYSOPHOSPHATIDYLCHOLINE L11046 S L1 AND ATHEROSCLEROSIS? L2L3 0 S L1 AND HYPERTWNSION? 39 S L2 AND HYPERTENSION? L4L5 1152 S L1 AND (CARDIOVASCULAR?) L6 304 S L2 AND L5 921 S L1 AND PHOSPHOCHOLINE? L7 2 S L6 AND L7 L8 2 DUPLICATE REMOVE L8 (0 DUPLICATES REMOVED) L9 9 S L5 AND L7 L10 9 DUPLICATE REMOVE L10 (0 DUPLICATES REMOVED) L11 L127 S L11 NOT L9 ·0 S L7 AND (METABOLIC SYNDROME) L130 S L13 AND HEART? L14

25 DUPLICATE REMOVE L15 (18 DUPLICATES REMOVED)

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L15

L16

L17

(FILE 'HOME' ENTERED AT 11:17:00 ON 28 FEB 2007)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 11:21:33 ON 28 FEB 2007

L1	20802 S LYSOPHOSPHATIDYLCHOLINE
<b>L</b> 2	1046 S L1 AND ATHEROSCLEROSIS?
L3	0 S L1 AND HYPERTWNSION?
L4	39 S L2 AND HYPERTENSION?
L5	1152 S L1 AND (CARDIOVASCULAR?)
L6	304 S L2 AND L5
L7	921 S L1 AND PHOSPHOCHOLINE?
L8	2 S L6 AND L7
L9	2 DUPLICATE REMOVE L8 (0 DUPLICATES REMOVED)
L10	9 S L5 AND L7
L11	9 DUPLICATE REMOVE L10 (0 DUPLICATES REMOVED)
L12	7 S L11 NOT L9
L13	0 S L7 AND (METABOLIC SYNDROME)
L14	0 S L13 AND HEART?
L15	43 S L7 AND HEART?
L16	25 DUPLICATE REMOVE L15 (18 DUPLICATES REMOVED)
L17	17 S L16 AND PD<1999

=>

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MEDLINE on STN
 ANSWER 14 OF 17
AN
     89322894
                  MEDLINE
DN
     PubMed ID: 2665794
TI
     Regulation of phosphatidylcholine metabolism in mammalian hearts
     Hatch G M; O K; Choy P C
ΑU
     Department of Biochemistry, Faculty of Medicine, University of Manitoba,
CS
     Winnipeg, Canada.
     Biochemistry and cell biology = Biochimie et biologie cellulaire,
SO
     (1989 Feb-Mar) Vol. 67, No. 2-3, pp. 67-77. Ref: 104
     Journal code: 8606068. ISSN: 0829-8211.
CY
     Canada
DT
     Journal; Article; (JOURNAL ARTICLE)
     (RESEARCH SUPPORT, NON-U.S. GOV'T)
     General Review; (REVIEW)
LA
     English
FS
     Priority Journals
EM
     198908
ED
     Entered STN: 9 Mar 1990
     Last Updated on STN: 9 Mar 1990
     Entered Medline: 31 Aug 1989
     Phosphatidylcholine is the major phospholipid in the mammalian
AB
     heart. Over 90% of the cardiac phosphatidylcholine is synthesized
     via the CDP-choline pathway. The rate-limiting step of this pathway is
     catalyzed by CTP:phosphocholine cytidylyltransferase. Current
     evidence suggests that phosphatidylcholine biosynthesis in the
    heart is regulated by the availability of CTP and the modulation
     of cytidylyltransferase activity. Phosphatidylcholine is degraded mainly
    by the actions of phospholipase A1 and A2, with the formation of
     lysophosphatidylcholine. Lysophosphatidylcholine may be
     further deacylated by lysophospholipase or reacylated back into the parent
    phospholipid by the action of acyltransferase. The accumulation of
     lysophosphatidylcholine in the heart may be one of the
    biochemical factors' for the production of cardiac arrhythmias.
CT
     Animals
       *Heart: PH, physiology
     *Mammals: ME, metabolism
     Mammals: PH, physiology
     *Myocardium: ME, metabolism
     *Phosphatidylcholines: ME, metabolism
```

Phosphatidylcholines: PH, physiology

0 (Phosphatidylcholines)

CN



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0829-8211



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biochimie et biologie cellulaire

Revue canadienne de biochimie et biologie cellulaire

Cover title: Canadian journal of biochemistry and cell biology =

Author: Canadian Biochemical Society.

> National Research Council Canada. Canadian Society for Cell Biology.

Ottawa: National Research Council of Canada = Conseil national des Imprint:

recherches du Canada, 1986-

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for Online version via Academic Search Premier (ASP). Feb

2001-

http://proquest.umi.com/pqdweb?

RQT=318&VName=PQD&clientid=19649&pmid=36120 Click here for Online version via Proguest. Jan 1, 1998-Present.

Notes:

URL:

Available on ADONIS, v. 73, no. 1-2 (1995) - v. 80, no. 4 (2002)

Includes bibliographies.

Articles in English; summaries in English and French.

Official journal of the Canadian Biochemical Society and the Canadian Society

for Cell Biology.

ISSN:

0829-8211

Subjects:

Biological chemistry -- Periodicals.

Cytology -- Periodicals.

Description: v.: ill.; 26 cm.

Continues: Canadian journal of biochemistry and cell biology

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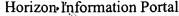
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ANSWER 1 OF 1
                   MEDLINE on STN
AN
     92197126
                  MEDLINE
DN
     PubMed ID: 1801455
     [Phospholipid thrombocyte activating factor, its analogs and antagonists:
ТT
     prospects of their use in medicine].
     Fosfolipidnyi faktor aktivatsii trombotsitov, ego analogi i antagonisty:
     perspektivy primeneniia v meditsine.
     Kulikov V I; Muzia G I
ΑU
     Vestnik Akademii meditsinskikh nauk SSSR, (1991) No. 10, pp. 13-7. Ref:
SO
     Journal code: 7506153. ISSN: 0002-3027.
CY
     USSR
DT
     (ENGLISH ABSTRACT)
     (IN VITRO)
     Journal; Article; (JOURNAL ARTICLE)
     General Review; (REVIEW)
LA
     Russian
     Priority Journals
FS
EM
     199204
ED
     Entered STN: 9 May 1992
     Last Updated on STN: 9 May 1992
     Entered Medline: 21 Apr 1992
     Experimental data on the biological activity of phospholipid
AΒ
     platelet-activation factor (PAF), its structural analogs and antagonists
     are discussed. The prospects of the use of PAF and PAF antagonists in
     medicine are under consideration. The conclusion is drawn that PAF
     antagonists may serve the basis for the development of highly potent drugs
     of new generation.
CT
      Azepines: DU, diagnostic use
     Azepines: PD, pharmacology
     *Azepines: TU, therapeutic use
     *Diterpenes
      Fibrinolytic Agents: PD, pharmacology
     *Fibrinolytic Agents: TU, therapeutic use
      Ginkgolides
      Humans
      Lactones: DU, diagnostic use
      Lactones: PD, pharmacology
     *Lactones: TU, therapeutic use
       *Lysophosphatidylcholines: PD, pharmacology
     *Platelet Activating Factor: AA, analogs & derivatives
      Platelet Activating Factor: AI, antagonists & inhibitors
     *Platelet Activating Factor: PH, physiology
      Platelet Activation: DE, drug effects
     *Platelet Activation: PH, physiology
      Platelet Aggregation: DE, drug effects
     *Platelet Aggregation: PH, physiology
      Platelet Function Tests
        Thrombosis: BL, blood
        Thrombosis: DT, drug therapy
       *Thrombosis: ET, etiology
      Triazoles: DU, diagnostic use
      Triazoles: PD, pharmacology
     *Triazoles: TU, therapeutic use
     105219-56-5 (WEB 2086); 99796-69-7 (ginkgolide B)
RN
     0 (1-acylglycerylphosphorylcholine); 0 (1-alkyl-2-acyl-sn-glycero-3-
CN
    phosphocholine); 0 (Azepines); 0 (Diterpenes); 0 (Fibrinolytic
    Agents); 0 (Ginkgolides); 0 (Lactones); 0 (Lysophosphatidylcholines**
       ); 0 (Platelet Activating Factor); 0 (Triazoles)
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```
ANSWER 2 OF 22 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN
AN
     1990:514666 BIOSIS
     PREV199090131942; BA90:131942
DN
     HYDROLYSIS OF 2 ACYL-SN-GLYCERO-3-PHOSPHOCHOLINES IN GUINEA-PIG
TI
     HEART MITOCHONDRIA.
     BADIANI K [Reprint author]; PAGE L; ARTHUR G
ΑU
     DEP BIOCHEM MOL BIOL, FAC MED, UNIV MANITOBA, 770 BANNATYNE AVE, MANIT,
CS
     CANADA R3E 0W3
     Biochemistry and Cell Biology, (1990) Vol. 68, No. 9, pp.
SO
     1090-1095.
     CODEN: BCBIEQ. ISSN: 0829-8211.
     Article
DT
FS
     BA
LΑ
     ENGLISH
     Entered STN: 19 Nov 1990
ED
     Last Updated on STN: 19 Nov 1990
     Although both 2-acyl-sn-glycero-3-phosphocholine and
AB
     1-acyl-sn-glycero-3-phosphocholine may be produced from
     phosphatidylcholine hydrolysis, studies on the former have lagged behind
     that of the latter. In this study a lysophospholipase A2 that hydrolyses
     2-acyl-sn-glycero-3-phosphocholine has been characterized in
     guinea pig heart mitochondria. The lysophospholipase A2 activity was not
     dependent on Ca2+ and was inhibited differentially by saturated and
     unsaturated fatty acids. This lysophospholipase A2 activity was able to
     discriminate among different molecular species of 2-acyl-sn-glycero-3-
     phosphocholines when they were presented individually or in pairs.
     The order of decreasing rates of hydrolysis of different molecular species
     of 2-lysophosphatidylcholines, when the substrates were
     presented singly, was 18:2 > 20:4 > 18:1 > 16:0. A differential
     inhibition of the rate of hydrolysis of the individual substrates was
     observed when the substrates were presented in pairs. The degree of
     inhibition was dependent on the molar ratio of the mixed substrates.
     characteristics of the enzyme suggest that involvement in the selective
     release of fatty acids from mitochondrial phosphatidylcholine would depend
     on a high selectivity of phospholipase A1 for different molecular species
     of phosphatidylcholine. A lysophospholipase A1 activity was also
     characterized in the mitochondria with a distinct acyl specificity from
     the lysophospholipase A2. Other characteristics of the two
     lysophospholipases suggest that the two reactions are not catalyzed by the
     same enzyme.
     Biochemistry studies - Proteins, peptides and amino acids
CC
                                                                 10064
     Biochemistry studies - Lipids
                                     10066
     Enzymes - Physiological studies
                                       10808
     Anatomy and Histology - Microscopic and ultramicroscopic anatomy
     Metabolism - Lipids
                           13006
     Cardiovascular system - Physiology and biochemistry 14504
IT
    Major Concepts
          Cardiovascular System (Transport and Circulation); Enzymology
        (Biochemistry and Molecular Biophysics); Metabolism; Morphology
IT
    Miscellaneous Descriptors
        FATTY ACID RELEASE
ORGN Classifier
                   86300
        Caviidae
     Super Taxa
       Rodentia; Mammalia; Vertebrata; Chordata; Animalia
       Animals, Chordates, Mammals, Nonhuman Vertebrates, Nonhuman Mammals,
```

Rodents, Vertebrates

```
ANSWER 2 OF 22 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN
     1990:514666 BIOSIS
AN
     PREV199090131942; BA90:131942
DN
     HYDROLYSIS OF 2 ACYL-SN-GLYCERO-3-PHOSPHOCHOLINES IN GUINEA-PIG
TI
     HEART MITOCHONDRIA.
     BADIANI K [Reprint author]; PAGE L; ARTHUR G
ΑU
     DEP BIOCHEM MOL BIOL, FAC MED, UNIV MANITOBA, 770 BANNATYNE AVE, MANIT,
CS
     CANADA R3E 0W3
     Biochemistry and Cell Biology, (1990) Vol. 68, No. 9, pp.
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     1090-1095.
     CODEN: BCBIEQ. ISSN: 0829-8211.
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     Article
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     BA
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     ENGLISH
ED
     Entered STN: 19 Nov 1990
     Last Updated on STN: 19 Nov 1990
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     Although both 2-acyl-sn-glycero-3-phosphocholine and
     1-acyl-sn-glycero-3-phosphocholine may be produced from
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CC
     Biochemistry studies - Proteins, peptides and amino acids
                                                                 10064
     Biochemistry studies - Lipids
                                     10066
     Enzymes - Physiological studies
                                       10808
     Anatomy and Histology - Microscopic and ultramicroscopic anatomy
                                                                        11108
    Metabolism - Lipids
                           13006
     Cardiovascular system - Physiology and biochemistry
                                                           14504
IT
    Major Concepts
          Cardiovascular System (Transport and Circulation); Enzymology
        (Biochemistry and Molecular Biophysics); Metabolism; Morphology
IT
    Miscellaneous Descriptors
        FATTY ACID RELEASE
ORGN Classifier
       Caviidae
                   86300
    Super Taxa
       Rodentia; Mammalia; Vertebrata; Chordata; Animalia
    Taxa Notes
       Animals, Chordates, Mammals, Nonhuman Vertebrates, Nonhuman Mammals,
```

Rodents, Vertebrates





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Revue canadienne de biochimie et biologie cellulaire

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Author: Canadian Biochemical Society.

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Imprint: Ottawa: National Research Council of Canada = Conseil national des

recherches du Canada, 1986-

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2001-

http://proquest.umi.com/pgdweb?

RQT=318&VName=PQD&clientid=19649&pmid=36120 Click here for Online version via Proquest. Jan 1, 1998-Present.

Notes:

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Cytology -- Periodicals.

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